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<b>Substitute for form 1449B/PTO</b>  <b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b>  (use as many sheets as necessary)		<b>Compleat if Known</b>			
		Application Number	09/853,753		
		Filing Date	May 14, 2001		
		First Named Inventor	B ch-Hansen		
		Group Art Unit	not assigned		
		Examiner Name	not assigned		
Sheet	1	of	1	Attorney Docket Number	45499.2

OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. <sup>1</sup>	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T <sup>2</sup>
OC	AA	Loss of Function Mutations in a Calcium-Channel $\alpha_1$ subunit gene in Xp11.23 cause incomplete X-linked congenital stationary night blindness - Paper by Bech-Hansen, et al. July, 1998.	
OC	AB	Mutations in NYX, encoding the leucine-rich proteoglycan nyctalopin, cause X-linked incomplete congenital stationary night blindness - Paper by Bech-Hansen et al. November, 2000.	
OC	AC	Evidence for Genetic Heterogeneity in X-linked Congenital Stationary Night Blindness - Paper by Bech-Hansen, et al. - published April 7, 1998.	
OC	AD	Leucine-Rich Repeat Glycoproteins of the Extracellular Matrix - Paper by Hocking, et al. accepted January 29, 1998.	

Examiner Signature	<i>Cheng Sun</i>	Date Considered	03/20/02
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